L1L2

L3

L4

L5 L6

L7

AN

AΒ

AB

(FILE 'HOME' ENTERED AT 09:56:11 ON 05 NOV 2002) FILE 'CA' ENTERED AT 09:56:21 ON 05 NOV 2002 E HAUTMAN D/AU 14 S E5-6 AND DRINK? 9 S L1 AND ION CHROMATOG? E JOYCE R/AU 46 S E7-9, E18-19, E23, E27 18 S L3 AND WATER 3 S L3 AND OXYHAL? 5 S L3 AND WATER/SO 16 S L2, L5-6

=> d 17 bib, ab 1-16

ANSWER 9 OF 16 CA COPYRIGHT 2002 ACS 118:109225 CA

TI Using ion chromatography to analyze inorganic disinfection by-products ΑU Hautman, Daniel P.; Bolyard, Michele

CS US Environ. Protect. Agency, Cincinnati, OH, 45268, USA

SO Journal - American Water Works Association (1992), 84(11), 88-93

Ion chromatog. is used to analyze drinking water for inorg. disinfection byproducts-the oxyhalides of Cl and Br. This investigation focused on stabilizing and preserving ClO2- by studying several agents known to quench its reaction with species present in drinking water. Based on the initial stability study, ethylenediamine was an effective preservative and was further studied using finished water from various utilities. Also, the measurement of BrO3- following ozonization of a river water matrix contq. 0.037 mg Br-/L is illustrated.

ANSWER 10 OF 16 CA COPYRIGHT 2002 ACS

117:118034 CA

Analysis of inorganic disinfection by-products using ion chromatography ΤI ΑU

Hautman, Daniel P.; Bolyard, Michele

CS Technol. Appl., Inc., Cincinnati, OH, 45268, USA

SO Proceedings - Water Quality Technology Conference (1992), Volume Date 1991, (Pt. 2), 1043-59

The US EPA is developing regulations for disinfection byproducts (DBPs). Ion chromatog. (IC) is used to analyze drinking water for inorg. DBPs that occur as the oxyhalides of Cl and Br. The oxyhalides of interest in this study are the anions ClO2-, ClO3-, and BrO3-. Since a previous study by J. D. Pfaff and C. A. Brockhoff (1990) found that ClO2- was unstable in drinking water, several species known to react with ClO2- and potentially present in **drinking** water were investigated in reagent-grade water. ability of different preservatives to stabilize ClO2- concns. in these reagent water matrixes was investigated. Based upon this initial stability study, ethylenediamine was found to be an effective preservative and was studied using finished drinking water from various utilities. Also, the formation of BrO3- following ozonization is illustrated with data from a pilot treatment plant.

- L7 ANSWER 11 OF 16 CA COPYRIGHT 2002 ACS
- AN 117:76098 CA
- TIAnalysis of oxyhalide disinfection by-products and other anions of interest in drinking water by ion chromatography
- ΑU Hautman, Daniel P.; Bolyard, Michele
- CS Technol. Appl. Inc., Cincinnati, OH, 45268, USA

SO Journal of Chromatography (1992), 602(1-2), 65-74

The US EPA is developing regulations for various drinking water disinfection byproducts (DBPs) that involves developing anal. methods for the DBPs formed as a result of different disinfection treatments and collecting occurrence data for these species. Ion chromatog. is one method being used to analyze drinking water samples for the following inorg. DBPs: Clo2-, Clo3-, and Bro3-. These anions, however, are difficult to sep. from the common interfering anions Cl-, Co32-, and No3-. A method is therefore presented by which tetraborate/boric acid is used to sep. these anions. Detection limits of the order of 10  $\mu$ g/L, using cond. and UV detection, were obtained. Stability studies of Clo2- showing the effectiveness of ethylenediamine as a preservative and summary data for an occurrence of nitrite, nitrate and DBP precursor bromide are presented.

L7 ANSWER 12 OF 16 CA COPYRIGHT 2002 ACS

AN 117:55526 CA

TI Occurrence of chlorate in hypochlorite solutions used for **drinking** water disinfection

AU Bolyard, Michele; Fair, Patricia Snyder; Hautman, Daniel P.

CS Tech. Support Div., U.S. Environ. Protect. Agency, Cincinnati, OH, 45268, USA

SO Environmental Science and Technology (1992), 26(8), 1663-5

AB / Hypochlorite solns. used for **drinking** water disinfection were obtained from 14 sites. In addn., source water samples (including both ground and surface waters) and **drinking** water samples were collected at these sites. Chlorate ion (ClO3-) concns. were detd. for all samples using **ion** chromatog. (10 μg/L reporting limit). Hypochlorite solns. contained 0.2-50

chromatog. (10  $\mu$ g/L reporting limit). Hypochlorite solns. contained 0.2-50 g/L ClO3-. Two of the source water samples had detectable levels of ClO3-, while all 14 **drinking** water samples contained ClO3- (ranging from 11 to 660  $\mu$ g/L). Hypochlorite solns. used to disinfect **drinking** water contain significant levels of ClO3-, and ClO3- is present in **drinking** water as a

direct result of this contamination.

ANSWER 13 OF 16 CA COPYRIGHT 2002 ACS

104:23835 CA

TI Ionic contamination: tracking it with ion chromatography

AU Joyce, R. J.

CS Dionex Corp., Sunnyvale, CA, 94088, USA

SO Ultrapure **Water** (1985), 2(4), 36-9

AB A review with 6 refs. on the development and state-of-the-art application of ion chromatog. for detn. of common anions and cations, carboxylates, and transition metal ions at ppb levels in industrial-use, high-purity waters is presented. Methods, including photometric detection and dual-column ion chromatog. in conjunction with effective cond. measurements, are emphasized.

=> log y STN INTERNATIONAL LOGOFF AT 10:04:22 ON 05 NOV 2002

=> d his

L1

L3

(FILE 'HOME' ENTERED AT 10:23:16 ON 05 NOV 2002)

FILE 'CA' ENTERED AT 10:23:27 ON 05 NOV 2002 10712 S (OXYHALIDE OR CLO2 OR CHLORINE DIOXIDE)

L2 1204 S L1(6A) (DETECT? OR DETERMIN? OR ASSAY? OR ANALY? OR ASSES? OR TEST?
OR MEASUR? OR MONITOR? OR ESTIMAT? OR EVALUAT? OR EXAMIN? OR SENSE# OR
SENSING OR SENSOR OR IDENTIF? OR PROBE# OR PROBING)

5 S L2 AND (BORIC OR BORATE OR BO3 OR HBO3)

```
191 S L1(5A) RESID?
L4
      309 S L2 AND (COLOR? OR DYE OR SPECTROPHOT? OR SPECTROMET? OR AZO? OR
L5
          PHOTOMET? OR REAGENT OR INDICATOR)
       20 S L4 AND L5
L6
      289 S L5 NOT L4
L7
L8
      202 S L7 AND (WATER OR H2O OR AQUEOUS OR STABILITY OR (CL2 OR CL OR
          CHLORINE) (5A) PRESEN? OR REAGENT OR INDICATOR)
L9
      226 S L3, L6, L8
L10
      204 S L9 NOT PY>1999
      171 S L10 NOT(STRATOS? OR CHEMAREG OR OSO4 OR RHODAMINE OR
L11
          PHENYLENEDIAMINE OR LIGNIN)
L12
      151 S L11 NOT (CHEMILUM? OR ENZYM? OR OXYGENAT? OR DIPOLE OR TRIPHEN? OR
          TRIS OR MULTISPEC?)
       20 S L11 NOT L12
L13
L14
        1 S L13 AND OXY CHLORINE
      116 S L12 NOT (METHYLENE BLUE OR PULP OR MASS OR KRAFT OR CHLOROPHENOL OR
L15
          TOLUIDINE)
       35 S L12 NOT L15
L16
L17
        2 S L16 AND (NH4OH OR METHYLORANGE)
       99 S L15 NOT(TROPOS? OR PHENYLBENZ? OR MICROB? OR TASTE OR ELECTRODE)
L18
       17 S L15 NOT L18
L19
L20
       1 S L19 AND NETWORK
L21
       98 S L18 NOT SPACE
```

## => d bib,ab 1-102 122

102 S L14, L17, L20-21

3 S L3 AND L22

L22 ANSWER 16 OF 102 CA COPYRIGHT 2002 ACS AN 123:237075 CA

TI Study of organic matter from raw and clarified waters by global analytical parameters

AU Lefebvre, E.

CS SAUR, Centre Pierre Crussard, Maurepas, 78310, Fr.

SO Revue des Sciences de l'Eau (1995), 8(1), 125-50

LA French

L22

L23

AΒ

The mol. wt. fractionation of dissolved org. matter in raw and clarified waters is described. Raw and clarified waters (resp. rw and cw) were sampled in SAUR water plants. All raw and clarified water samples were characterized according to total org. carbon (TOC), UV absorbance (254 nm), and trihalomethane formation potential (THMFP) under the following conditions: about 20°C, 4 mg Cl2/mg TOC, and a 72-h contact time in the dark. A Dohrmann DC80 and a Uvikon 930 were used for the detn. of TOC (DOC) and UV-absorbance at 254 nm, resp. When a preoxydation step was employed at the water plant, the clarification treatment was performed with a lab. app. Bioeliminable Org. Dissolved Carbon in water was detd. by the method described. Cl2 and ClO2 demands of raw and clarified waters were conducted as batch operations with oxidant doses of 1, 2, and 4 mg oxidant per mg TOC. Residual chlorine and chlorine dioxide in solns. were detd. resp. by spectrophotometric measurements by two colorimetric DPD and ACVK methods.

- L22 ANSWER 23 OF 102 CA COPYRIGHT 2002 ACS
- AN 119:194616 CA
- TI Effect of dimethyl sulfoxide as a masking agent for chlorine in the selective **determination** of **aqueous chlorine dioxide**
- AU Imaizumi, Noriko; Nakahara, Yumiko; Suzuki, Kayoko; Oikawa, Kikuo
- CS Niigata Coll. Pharm., Niigata, 950-21, Japan

Chemistry Letters (1993), (8), 1333-6

AB
For the selective detn. of aq. chlorine dioxide (C102) in the mixed soln. with chlorine, DMSO (DMSO) was used as a masking agent for chlorine. In spectrophotometric and iodometric detn. of C102, a large excess of DMSO did not have an effect on C102, but it completely depressed the interference from chlorine.

L22 ANSWER 31 OF 102 CA COPYRIGHT 2002 ACS

AN 115:120902 CA

TI Computer optimization in ion chromatography. II. A systematic evaluation of linear retention models for anions

AU Sosimenko, Andrew D.; Haddad, Paul R.

CS Dep. Anal. Chem., Univ. New South Wales, Kensington, 2033, Australia

SO Journal of Chromatography (1991), 546(1-2), 37-59

Extensive retention data for non-suppressed ion chromatog. of anions were ABacquired for 17 analytes (halides, oxyhalides, nitrite, nitrate, sulfite, sulfate, bisulfite, thiosulfate, phosphate, thiocyanate, carbonate, acetate, and oxalate) on 3 stationary phases (Waters IC Pak A, Hamilton PRP-X100 and Vydac 302.IC 4.6) by using 7 eluent types (benzoate, phthalate, hydroxide, carbonate/bicarbonate, gluconate/borate, ptoluenesulfonate, and phosphate). These retention data are used to assess the validity of retention models which predict a linear relation between the logarithm of solute capacity factor and the logarithm of the activity of the eluent competing anion. The linearity of these plots is uniformly good, but the slopes differ markedly from those predicted from theory. When the eluent contains 2 competing anions, neither the dominant equil. approach nor the effective charge approach give reliable prediction of the slopes. Optimization of one eluent parameter at a time (e.g., the concn. of the competing anion in the eluent) can be successful if the slope of the retention plot is detd. by measurement of analyte retention times at 2 eluent compns: falling at the extremes of the range of eluent compns. under consideration. An example of this "end points" method is provided, in which the concn. of a phthalate eluent is optimized.

ANSWER 39 OF 102 CA COPYRIGHT 2002 ACS 109:196775 CA

Selection of an **analytical** technique to **measure chlorine dioxide** in the field and **determination** of residual effectiveness

AU Thompson, Artis; Matthews, Nancy; Myers, Gordon L.; Owen, Douglas M.

CS Galveston Cty. Water Auth., Texas City, TX, USA

SO Proceedings - Water Quality Technology Conference (1988), Volume Date 1987, 15(Issue Answers Today's Water Qual. Prof.), 1043-54

colorimetric methods if consistent and reproducible procedures are followed. Samples collected in the field must be shielded from light prior to anal. to avoid photolytic decompn. A sample bomb can be used effectively to collect and store samples for anal. Samples collected in the field should be analyzed immediately to avoid possible decompositional effects during transport. Bacterial counts after introducing ClO2 as a codisinfectant may be high initially as a result of stripping of corrosion byproducts and other depositional layers on the inside of distribution piping, in which bacteria reside. Bacterial counts decrease with time as these materials are flushed from the system. Only white-staining gram pos. rods show resistance to the oxidizing effects of ClO2 in the distribution system.

L22

AΒ

ANSWER 40 OF 102 CA COPYRIGHT 2002 ACS 109:196562 CA

- TI A critical review of the analytical methods currently used for the measurement of free, combined, and **oxy-chlorine** species
- AU Gordon, Gilbert; Pacey, Gilbert E.; Cooper, William J.
- CS Dep. Chem., Miami Univ., Oxford, OH, 45056, USA
- SO Proceedings Water Quality Technology Conference (1988), Volume Date 1987, 15(Issue Answers Today's Water Qual. Prof.), 1005-42
- The review and discussion, with 75 refs., covers the measurement of free-, combined, and oxy-Cl species in water, including Cl and chloramine chem., ClO2 chem., potential interferences, Cl conversions, UV spectrometric detn. of Cl and chloramine, amperometric and iodometric titrns., colorimetric methods, electrode methods, chemiluminescence method, ClO2- and ClO3-detn., and flow injection anal.
- L22 ANSWER 47 OF 102 CA COPYRIGHT 2002 ACS
- AN 103:188693 CA
- TI Selective **determination** of **chlorine dioxide** using gas diffusion flow injection **analysis**
- AU Hollowell, David A.; Pacey, Gilbert E.; Gordon, Gilbert
- CS Dep. Chem., Miami Univ., Oxford, OH, 45056, USA
- SO Anal. Chem. (1985), 57(14), 2851-4
- AB An automated absorbance technique for the detn. of aq. ClO2 was developed by using gas diffusion flow injection anal. A gas diffusion membrane is used to sep. the donor (sampling) stream from the acceptor (detecting) stream. The absorbance of ClO2 was monitored at 359 nm. The first method uses H2O as the acceptor stream and gives a detection limit of 0.25 mg ClO2/L. This system is >550 times more selective for ClO2 than Cl. To further minimize Cl interference, oxalic acid is used in the acceptor stream. The detection limit for this system is 0.45 mg ClO2/L. This second system is >5400 times more selective for ClO2 than Cl. Both methods show excellent selectivity for ClO2 over Fe and Mn compds., as well as other oxychlorinated compds. such as ClO3- and ClO4-.



ANSWER 59 OF 102 CA COPYRIGHT 2002 ACS

93:191731 CA

- TI Comparison between **colorimetric** and electrometric methods for chlorine and its derivative compounds
- AU Piccardi, Giovanni; Barbolani, Emilia; Pantani, Francesco
- CS Inst. Anal. Chem., Univ. Florence, Florence, 50121, Italy
- SO Water, Air, Soil Pollut. (1980), 13(2), 197-205
- AB A crit. comparison of the usual methods of detg. Cl and its compds., particularly those used as sterilants in potable water plants, showed that the o-tolidine test is neither reproducible nor specific. ClO2 is best detd. by Acid Chrome Violet K. The role of amperometric titrn. in distinguishing between the various sterilants and in examg. the reactions involved in water chlorination is discussed.



ANSWER 78 OF 102 CA COPYRIGHT 2002 ACS

70:117967 CA

- TI Behavior and determination of chlorine dioxide
- AU Myhrstad, Jan A.; Samdal, J. E.
- CS Norw. Inst. Water Res., Oslo, Norway
- SO J. Amer. Water Works Ass. (1969), 61(4), 205-8
- AB A new combination of methods for detg. ClO2, ClO2-, and Cl2 in water is discussed. The method is based on direct photometric detn. of ClO2 with Acid Chrome Violet K; iodometric-potentiometric titrn. at pH 7 to give Cl2 + 0.80 ClO2; and an iodometric-potentiometric titrn. at pH 7 after acidifying to pH 2.5-3, to give Cl2 + ClO2 + ClO2-.

122

ANSWER 89 OF 102 CA COPYRIGHT 2002 ACS

AN 59:67991 CA OREF 59:12507a-b

TI The colorimetric determination of chlorine dioxide in the presence of chlorine in water

AU Merenyi, P.; Kuba, P.

CS & Chem. Zavody J. Dimitrova, Bratislava, Czech.

SO Chem. Zvesti (1963), 17, 146-51

AB By applying ClO2 for water treatment it is necessary to det. its residual content in the treated water in the presence of Cl. The use of tyrosine for this purpose is discussed. To eliminate the interfering effect of Cl, monoethylamine is applied.

122 AN

ANSWER 92 OF 102 CA COPYRIGHT 2002 ACS

AN 55:53203 CA OREF 55:10191g-h

TI Determination of chlorine dioxide in concentrated solutions in the presence of chlorine

AU 'Lepeintre, Marcel; Dupuy, Jeanine; Ouvard, Jean

SO & Chim. anal. (1960), 42, 498-500

AB ClO2 forms a colored product with tyrosine. The interference of Cl is removed by fixation with EtNH2. The absorption of the colored product is measured at 490 m $\mu$  with an error within 5%.

=> log y STN INTERNATIONAL LOGOFF AT 11:46:59 ON 05 NOV 2002 From: Sent: 🌌 Mellerson, Kendra

Tuesday, November 05, 2002 2:26 PM

To: Subject:

STIC-ILL FW: ill request

----Original Message----

Fr m:

Soderquist, Arlen

Sent:

Tuesday, November 05, 2002 2:02 PM

T:

STIC-EIC1700

Subject:

ill request

Arlen Soderquist

AU 1743

308-3989

CP3-7A11

Serial No. 09/394647

Needed by 11-10-02

Abstract

L7 ANSWER 10 OF 16 CA COPYRIGHT 2002 ACS

AN 117:118034 CA

TI Analysis of inorganic disinfection by-products using ion chromatography

AU Hautman, Daniel P.; Bolyard, Michele

CS Technol. Appl., Inc., Cincinnati, OH, 45268, USA

SO Proceedings - Water Quality Technology Conference (1992), Volume Date 1991, (Pt. 2), 1043-59

AB The US EPA is developing regulations for disinfection byproducts (DBPs). Ion chromatog. (IC) is used to analyze drinking water for inorg. DBPs that occur as the oxyhalides of Cl and Br. The oxyhalides of interest in this study are the anions ClO2-, ClO3-, and BrO3-. Since a previous study by J. D. Pfaff and C. A. Brockhoff (1990) found that ClO2- was unstable in drinking water, several species known to react with ClO2- and potentially present in drinking water were investigated in reagent-grade water. The ability of different preservatives to stabilize ClO2- concns. in these reagent water matrixes was investigated. Based upon this initial stability study, ethylenediamine was found to be an effective preservative and was studied using finished drinking water from various utilities. Also, the formation of BrO3- following ozonization is illustrated with data from a pilot treatment plant.



From:

Mellerson, Kendra

Sent:

Tuesday, November 05, 2002 2:26 PM

To: Subject:

STIC-ILL FW: ill request

----Original Message----

Fr m:

Soderquist, Arlen

Sent:

Tuesday, November 05, 2002 1:56 PM

T :

STIC-EIC1700

Subject:

ill request

Arlen Soderquist

AU 1743

308-3989

CP3-7A11

Serial No. 09/394647

Needed by 11-10-02

Abstract

L22 ANSWER 39 OF 102 CA COPYRIGHT 2002 ACS

AN109:196775 CA

Selection of an analytical technique to measure chlorine dioxide in the TI field and determination of residual effectiveness

Thompson, Artis; Matthews, Nancy; Myers, Gordon L.; Owen, Douglas M. ΑU

Galveston Cty. Water Auth., Texas City, TX, USA CS

Proceedings - Water Quality Technology Conference (1988), Volume Date SO 1987, 15(Issue Answers Today's Water Qual. Prof.), 1043-54

Clo2 residuals can be measured reliably in water distribution systems AB using colorimetric methods if consistent and reproducible procedures are Samples collected in the field must be shielded from light prior to anal. to avoid photolytic decompn. A sample bomb can be used effectively to collect and store samples for anal. Samples collected in the field should be analyzed immediately to avoid possible decompositional effects during transport. Bacterial counts after introducing ClO2 as a codisinfectant may be high initially as a result of stripping of corrosion byproducts and other depositional layers on the inside of distribution piping, in which bacteria reside. Bacterial counts decrease with time as these materials are flushed from the system. Only white-staining gram pos. rods show resistance to the oxidizing effects of ClO2 in the distribution system.

CONPLETE

From: <sup>a</sup> Sent: To: Subject:

Mellerson, Kendra Tuesday, November 05, 2002 2:26 PM STIC-ILL FW: ill request

----Original Message-----

From:

Soderquist, Arlen

Sent:

Tuesday, November 05, 2002 1:55 PM

To:

STIC-EIC1700

Subject:

ill request

Arlen Soderquist

AU 1743

308-3989

CP3-7A11

Serial No. 09/394647

Needed by 11-10-02

Abstract

L22 ANSWER 40 OF 102 CA COPYRIGHT 2002 ACS

AN 109:196562 CA

A critical review of the analytical methods currently used for the measurement of TI free, combined, and oxy-chlorine species

Gordon, Gilbert; Pacey, Gilbert E.; Cooper, William J. ΑU

CS Dep. Chem., Miami Univ., Oxford, OH, 45056, USA

SO Proceedings - Water Quality Technology Conference (1988), Volume Date 1987, 15(Issue Answers Today's Water Qual. Prof.), 1005-42

The review and discussion, with 75 refs., covers the measurement of AB free-, combined, and oxy-Cl species in water, including Cl and chloramine chem., ClO2 chem., potential interferences, Cl conversions, UV spectrometric detn. of Cl and chloramine, amperometric and iodometric titrns., colorimetric methods, electrode methods, chemiluminescence method, ClO2- and ClO3- detn., and flow injection anal.

From:

Mellerson, Kendra

Sent:

Tuesday, November 05, 2002 2:26 PM

To: Subject:

STIC-ILL FW: ill request

----Original Message-----

Fr m:

Soderquist, Arlen

Sent:

Tuesday, November 05, 2002 1:51 PM

To:

STIC-EIC1700

Subject:

ill request

Arlen Soderquist

AU 1743

308-3989

CP3-7A11

Serial No. 09/394647

Needed by 11-10-02

Abstract

L22 ANSWER 89 OF 102 CA COPYRIGHT 2002 ACS

59:67991 CA AN

OREF 59:12507a-b

The colorimetric determination of chlorine dioxide in the presence of TI chlorine in water

ΑU Kerenyi, P.; Kuba, P.

Chem. Zavody J. Dimitrova, Bratislava, Czech. CS

Chem. Zvesti (1963), 17, 146-51 SO

AB By applying ClO2 for water treatment it is necessary to det. its residual content in the treated water in the presence of Cl. tyrosine for this purpose is discussed. To eliminate the interfering effect of Cl, monoethylamine is applied.

From:

Mellerson, Kendra

Sent: To:

Tuesday, November 05, 2002 2:26 PM

Subject:

STIC-ILL FW: ill request

----Original Message----

Fr m:

Soderquist, Arlen

Sent: .

Tuesday, November 05, 2002 1:52 PM

Т:

STIC-EIC1700

Subject:

ill request

Arlen Soderquist AU 1743

308-3989

CP3-7A11

Serial No. 09/394647

Needed by 11-10-02

Abstract

L22 ANSWER 78 OF 102 CA COPYRIGHT 2002 ACS

AN 70:117967 CA

TI Behavior and determination of chlorine dioxide

ΑU

Myhrstad, Jan A.; Samdal, J. E.

CS Norw. Inst. Water Res., Oslo, Norway

SO J. Amer. Water Works Ass. (1969), 61(4), 205-8

A new combination of methods for detg. ClO2, ClO2-, and Cl2 in water AB The method is based on direct photometric detn. of ClO2 with is discussed. Acid Chrome Violet K; iodometric-potentiometric titrn. at pH 7 to give Cl2 + 0.80 ClO2; and an iodometric-potentiometric titrn. at pH 7 after acidifying to pH 2.5-3, to give Cl2 + Cl02 + Cl02-.

From: Sent:

Mellerson, Kendra

Tuesday, November 05, 2002 2:26 PM

To: Subjec /

STIC-ILL FW: ill request

----Original Message----

Fr m:

Soderquist, Arlen

Sent:

Tuesday, November 05, 2002 1:53 PM

T :

STIC-EIC1700

Subject:

ill request

Arlen Soderquist

AU 1743

308-3989

CP3-7A11

Serial No. 09/394647

Needed by 11-10-02

Abstract

L22 ANSWER 59 OF 102 CA COPYRIGHT 2002 ACS

ΑN 93:191731 CA

Comparison between colorimetric and electrometric methods for chlorine TIand its derivative compounds

ΑU Piccardi, Giovanni; Barbolani, Emilia; Pantani, Francesco

CS Inst. Anal. Chem., Univ. Florence, Florence, 50121, Italy

Water, Air, Soil Pollut. (1980), 13(2), 197-205 SO

A crit. comparison of the usual methods of detg. Cl and its compds., particularly those used as sterilants in potable water plants, showed that the o-tolidine test is neither reproducible nor specific. Clo2 is best detd. by Acid Chrome Violet K. The role of amperometric titrn. in distinguishing between the various sterilants and in examg. the reactions involved in water chlorination is discussed.

From:

Mellerson, Kendra

Sent:

Tuesday, November 05, 2002 2:26 PM

To: Subject?

STIC-ILL FW: ill request

----Original Message----

From:

Soderquist, Arlen

Sent:

Tuesday, November 05, 2002 1:59 PM

To:

STIC-EIC1700

Subject:

ill request

Arlen Soderquist

AU 1743

308-3989

CP3-7A11

Serial No. 09/394647

Needed by 11-10-02

Abstract

L22

ANSWER 16 OF 102 CA COPYRIGHT 2002 ACS

AN 123:237075 CA

Study of organic matter from raw and clarified waters by global analytical TI parameters

ΑU Lefebvre, E.

SAUR, Centre Pierre Crussard, Maurepas, 78310, Fr. CS

SO Revue des Sciences de l'Eau (1995), 8(1), 125-50

LA

The mol. wt. fractionation of dissolved org. matter in raw and clarified waters is AB described. Raw and clarified waters (resp. rw and cw) were sampled in SAUR water plants. All raw and clarified water samples were characterized according to total org. carbon (TOC), UV absorbance (254 nm), and trihalomethane formation potential (THMFP) under the following conditions: about 20 C, 4 mg Cl2/mg TOC, and a 72-h contact time in the dark. Dohrmann DC80 and a Uvikon 930 were used for the detn. of TOC (DOC) and UV-absorbance at 254 nm, resp. When a preoxydation step was employed at the water plant, the clarification treatment was performed with a lab. app. Bioeliminable Org. Dissolved Carbon in water was detd. by the method described. Cl2 and ClO2 demands of raw and clarified waters were conducted as batch operations with oxidant doses of 1, 2, and 4 mg oxidant per mg TOC. Residual chlorine and chlorine dioxide in solns. were detd. resp. by spectrophotometric measurements by two colorimetric DPD and ACVK methods.

'MMPLETED

From:

Mellerson, Kendra

Sent:

Tuesday, November 05, 2002 2:26 PM

To: Subject: STIC-ILL FW: ill request

----Original Message----

Fr m:

Soderquist, Arlen

Sent:

Tuesday, November 05, 2002 2:03 PM

Т:

STIC-EIC1700

Subject:

ill request

Arlen Soderquist

AU 1743

308-3989

CP3-7A11

Serial No. 09/394647

Needed by 11-10-02

Abstract

L7

ANSWER 9 OF 16 CA COPYRIGHT 2002 ACS

ΑN 118:109225 CA

Using ion chromatography to analyze inorganic disinfection by-products TI

ΑU Hautman, Daniel P.; Bolyard, Michele

CS US Environ. Protect. Agency, Cincinnati, OH, 45268, USA

SO Journal - American Water Works Association (1992), 84(11), 88-93

Ion chromatog. is used to analyze drinking water for inorg. AB

disinfection byproducts-the oxyhalides of Cl and Br. This investigation focused on stabilizing and preserving ClO2- by studying several agents known to quench its reaction with species present in drinking water. Based on the initial stability study, ethylenediamine was an effective preservative and was further studied using finished water from various utilities. Also, the measurement of BrO3- following ozonization of a river water matrix contg. 0.037 mg Br-/L is illustrated.

From: Sent: To: Subject:

Mellerson, Kendra 🖈 Tuesday, November 05, 2002 2:26 PM STIC-ILL FW: ill request

----Original Message----

From:

Soderquist, Arlen

Sent:

Tuesday, November 05, 2002 1:49 PM

T:

STIC-EIC1700

Subject:

ill request

Arlen Soderquist

AU 1743

308-3989

CP3-7A11

Serial No. 09/394647

Needed by 11-10-02

Abstract

OREF 55:10191g-h

L22 ANSWER 92 OF 102 CA COPYRIGHT 2002 ACS AN 55:53203 CA

Determination of chlorine dioxide in concentrated solutions in the presence of chlorine

Lepeintre, Marcel; Dupuy, Jeanine; Ouvard, Jean ΑU

SO Chim. anal. (1960), 42, 498-500

AΒ ClO2 forms a colored product with tyrosine. The interference of Cl is removed by fixation with EtNH2. The absorption of the colored product is measured at 490 m $\mu$  with an error within 5%.



From: Sent:

Mellerson, Kendra

To: Subject: Tuesday, November 05, 2002 2:26 PM

STIC-ILL FW: ill request

----Original Message----

From:

Soderquist, Arlen

Sent:

Tuesday, November 05, 2002 2:00 PM

T:

STIC-EIC1700

Subject:

ill request

Arlen Soderquist

AU 1743

308-3989

CP3-7A11

Serial No. 09/394647

Needed by 11-10-02

Abstract

L7

ANSWER 13 OF 16 CA COPYRIGHT 2002 ACS

AN 104:23835 CA

ΤI Ionic contamination: tracking it with ion chromatography

ΑU Joyce, R. J.

CS Dionex Corp., Sunnyvale, CA, 94088, USA

SO Ultrapure Water (1985), 2(4), 36-9

A review with 6 refs. on the development and state-of-the-art AB application of ion chromatog. for detn. of common anions and cations, carboxylates, and transition metal ions at ppb levels in industrial-use, high-purity waters is presented. Methods, including photometric detection and dual-column ion chromatog. in conjunction with effective cond. measurements, are emphasized.